

RESULTS OF FAST $O(^3P)$ COLLISIONS WITH C_2H_2 , PROGRESS ON $O(^3P)$ EXCITATION OF $HNCO$, AND DETECTION OF NEW DISSOCIATIVE CHANNELS IN NO

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Using the JPL atomic-oxygen facility, we have observed optical emission in gas-phase single collisions of fast (3-22 eV translational energy) $O(^3P)$ atoms with acetylene. Optical emissions were measured in the wavelength range 300-850 nm, and corresponded to the $CH A^2\Delta \rightarrow X^211$, and $B^2\Sigma^- \rightarrow X^211$, transitions [1]. The $A \rightarrow X$ and $B \rightarrow X$ emissions were fit to a synthetic spectrum of $CH(A)$ at a vibrational temperature T_v of 10000 K (0.86 eV) and a rotational temperature T_r of approximately 5000 K (0.43 eV); and $CH(B)$ to $T_v = 2500$ K (0.22 eV) and $T_r = 1\,000$ K (0.09 eV). The energy threshold for the $A \rightarrow X$ emission was measured to be 7.3 ± 0.4 eV (lab) or 4.5 ± 0.2 eV (c. m.). This agrees with the energy threshold of 7.36 eV (lab) for the reaction $O(^3P) + C_2H_2 \rightarrow CH(A) + HCO$.

Work in progress involves the synthesis of the unstable molecule $HNCO$. The simplest method appears to be by dropwise addition of a concentrated solution of potassium cyanate to phosphoric acid, with collection of the evolved $HNCO$ in a dry-ice acetone trap. The excitation spectrum of $O(^3P) + HNCO$ will be studied, and correlations made with shuttle-exhaust observations in several recent flights.

Finally recent results, using the same experimental apparatus, of dissociative electron attachment to NO will be presented [2]. In addition to the known $N(^2D^o) + O^-(^2P)$ channel, two new channels $N(^4S^o) + O^-(^2P)$ and $N(^2P^o) + O^-(^2P)$ were detected. Cross sections for each of the channels were obtained by normalizing the scattering intensities to previously-measured total cross sections.

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[1] O. J. Orient, A. Chutjian, and E. Murad, *Phys. Rev. A* 57, 2094 (1995).

[2] O. J. Orient and A. Chutjian, *Phys. Rev. Letters* 74, 5017 (1995).